## **AMENDMENTS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

	In the Claims
	Claims 1-5 (cancelled)
H Sydney Commencer	Claim (currently amended): An isolated compound comprising a contiguous sequence of amino acids within the sequence representing residues 149-197177 of the G protein of respiratory syncytial virus (RSV), wherein more than one of cysteines 173, and 176, 182 and 186
a gr	is are absent or blocked, wherein said compound is not glycosylated, and wherein said compound human has the ability to inhibit infectivity of RSV.
	Claims 7-8 (cancelled)
# G	Claim (previously amended): A compound according to claim 6, wherein one or more amino acids is replaced by its corresponding D-amino acid.
Swit 2	Claim 10 (cancelled)
# 2	Claim It (previously amended): A compound according to claim to, wherein the compound is labelled with a detectable marker.
*	Claim 12 (previously amended): A compound according to claim 11, wherein the detectable marker is a radioactive label.
H	Claim 13 (previously amended): A compound according to claim 11, wherein the detectable marker is a fluorescent, chemiluminescent or enzymic marker.

Claims 14-15 (cancelled)

Claim 16 (withdrawn): An antibody directed against a compound selected from the group consisting of the compounds of Claims 1 to 10.

Claim 17 (withdrawn): An antibody according to Claim 16 which is a protective antibody.

Claim 18 (withdrawn): A composition comprising antibody selected from the group of the antibodies of Claim 16 and Claim 17.

Claims 19-20 (cancelled)

Claim 21 (withdrawn): A composition according to any one of Claim 16 in which the virus is human RSV.

Claim 22 (cancelled)

Claim 23 (withdrawn): A method of diagnosis of *Pneumovirus* infection, comprising exposing a biological fluid or sample from a mammal suspected of being infected with said virus to a compound selected from the group consisting of the compounds of Claims 1 to 10, and measuring the interaction between the compound and said fluid or sample.

Claim 24 (cancelled)

Claim 25 (withdrawn): A method of identification of a cell surface receptor for respiratory syncytial virus G protein, comprising the step of detection of binding of a compound selected from the group consisting of the compounds of Claims 11 to 13 to a cell surface protein.

Claim 26 (cancelled)

Claim 27 (withdrawh): A method according to Claim 25, in which the cell is susceptible to infection by respiratory syncytial virus.

Claim 28 (withdrawn): A method according to Claim 25, in which the cell is a HEp-2 cell.

Claim 29 (withdrawn): A method according to Claim 25, in which the method comprises the step of photoaffinity labelling of the receptor with a benzoylbenzyl derivative of the compound.

Claim 30 (withdrawn): A method according to Claim 25, in which the method comprises the step of labelling of the receptor with a fluorescent derivative of the compound.

Claim 31 (withdrawn): A method according to Claim 25, in which the method comprises the steps of binding a biotinylated derivative of the compound to a receptor, and binding of avidin to the derivative.

Claim 32 (withdrawn): A method according to Claim 25, in which the method comprises the step of using an antibody according to Claim 16 to detect the binding of the compound.

Claim 33 (withdrawn): A method according to Claim 25, in which the compound is multiply derivatised, thereby to achieve combined cross-linking, detection and identification of a receptor.

Claim 34 (previously amended): A compound according to claim 8, wherein the contiguous sequence represents residues 149 to 177 of the G protein of RSV.

Claim 35 (previously amended): A diagnostic composition comprising a compound according to claim 6.

Claim 36 (cancelled)

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Serial No. 09/202,035 Docket No. 273402004000 Claim 3 (previously added): A diagnostic composition according to claim 36, wherein one or more amino acids is replaced by its corresponding D-amino acid.

Claim 38 (cancelled)

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Claim 39 (previously amended): A composition comprising a compound according to claim 4, together with a pharmaceutically acceptable carrier.

Claim 40 (cancelled)

Claim of (previously amended): A composition comprising a compound according to claim, wherein one or more amino acids is replaced by its corresponding D-amino acid.

Claim 42 (cancelled)

Claim 33 (previously added): Accompound according to claim 5, comprising the amino acid sequence KQRQNKPPSKPNNDFHFEVFNFVPCSICG (SEQ ID NO:39), wherein the cysteine residues are derivatized with acetamidomethyl.

Claim 44 (previously added): A compound according to claim 6, consisting of acetyl-KQRQNKPPSKPNNDFHFEVFNFVPCSICGAmide (SEQ ID NO:39), wherein the cysteine residues are derivatized with acetamidomethyl.

Claim 45 (previously added): A method of inhibiting the cytopathic effect of RSV, polynerical comprising contacting an RSV susceptible cell with the compound of claim 5.

Claim 46 (previously added): A method according to claim 45, wherein the contiguous huma u sequence of amino acids represents residues 149 to 177 of the G protein of RSV.

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Claim 47 (previously added): A method according to claim 45, wherein the compound comprises the amino acid sequence KQRQNKPPSKPNNDFHFEVFNFVPCSICG (SEQ ID NO:39), wherein the cysteine residues are derivatized with acetamidomethyl.

Claim (previously added): A method according to claim 45, wherein the compound is acetyl-KQRQNKPPSKPNNDFHFEVFNFVPCSICGAmide (SEQ ID NO:39), wherein the cysteine residues are derivatized with acetamidomethyl.

Claim 49 (previously added): A method of inhibiting the cytopathic effect of RSV, comprising contacting an RSV susceptible cell with a compound comprising a contiguous sequence of amino acids within the sequence representing residues 149-197 of the G protein of respiratory syncytial virus (RSV).

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Claim 10 (currently amended): A compound comprising a contiguous sequence of amino acids within the sequence representing residues 149-197177 of the G protein of respiratory syncytial virus (RSV), wherein none neither of cysteines 173; or 176, 182 and 186 is functional to polynerial form a disulfide bridge, wherein said compound is not glycosylated, and wherein said compound has the ability to inhibit infectivity of RSV.